PTO/SB/08s (08-03.)
Approved for use through 07/31/2006, OMB 0651-0031
U.S. Patient and Trademark Office, U.S. DEPARTMENT OF COMMERCE

US Research use brough US Reduction Act of 1995, no persones are required to respect to a collection of information unites of contains a visit OMS control number.

Application Number

Filing Date

INCODMATION DISCLOSURE

STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)					First Named Inventor Maurice Howard Fisher						
					Art Unit						
					Examiner Name			-			
					Attorney Docket Num			1010-00400			
					U.S.I	PATENTS				Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue E	e Date Name of Patentee of Applicant R		Releva	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear			
	1										
If you wish	h to a	dd additional U.S. Pater	nt citatio	n inform	ation pl	ease click	the A	dd button.		Add	
			U.S.P	ATENT	ATENT APPLICATION PUBLICATIONS					Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹		to Name of Patentee or Applicant R		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear				
	1	20020017642		2002-0	2-14	Mizushima Kazuki et al.					
	2	20020125475		2002-0	9-12	Chu Jack Oon et al.					
If you wish	h to a	dd additional U.S. Publi	shed Ap	plication	citation	n informati	on ple	ase click the Ad	d button	Add	
				FOREIG	SN PAT	ENT DOC	UME	NTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³			Kind Code ⁴	Publication Date	on	Applicant of cited		Pages,Columns,Lir where Relevant Passages or Relev rigures Appear	76
	1	2003 197544	JP			2003-11-0	15	Sumitomo Mitsubis	ihi		
	2	03/103031	wo			2003-12-1	11	University of Warw	ick		

Application Number | Filing Date | Find The Tender | Find The Tend

If you wish	n to a	dd add	ditional Foreign Patent Document citation information please cl	ick the Add button	Add	
			NON-PATENT LITERATURE DOCUMEN	ITS	Remove	
Examiner Initials*	Cite No	(bool	de name of the author (in CAPITAL LETTERS), title of the arti- k, magazine, journal, serial, symposium, catalog, etc), date, pa isher, city and/or country where published.			Ţs
	1		ACCHI ET AL., Continuously graded buffers for InGaAs/GaAs structu wth 175/176 (1997) 1009-1015	res grown on GaAs	; Journal of Crystal	
	2	PCT	International Search Report for International Application PCT/GB200	5/000490, dated Jul	y 6, 2005 (4 p.)	
If you wish	h to a	dd add	ditional non-patent literature document citation information plea	se click the Add b	utton Add	
			EXAMINER SIGNATURE			
Examiner	Signa	ture	Di	ate Considered		

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

See for Codes of USPTO Platent Documents at year, USPTO, DOC or MPEP 901.04. * Enter office and assess the document, by the to-elect code (VIPO Standard STI), 3.** Explaines parted to convent, by an advanced strain and the Explaines special convenient, be noticated on the parent for superior the Emperor must proceed the senior number for papent document. *
**Indicate of coursent by the appropriate symbols as indicated on the document under WIPO Standard STI, 16 if possible. *
Applicant is to place a check mark here fittingle language strated inst attacked.

Application Number Fling Date Fling Date STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99) Application Number Fline Fli

CERTIFICATION STATEMENT

Please see	37	CFR :	1.97	and 1	1.98 to make	the appro	priate	selection	(s):	
------------	----	-------	------	-------	--------------	-----------	--------	-----------	------	--

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 3.7 CFR 197(eVI).

ΩR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(s)(c).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- .7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/ttumey/	Date (YYYY-MM-DD)	2006-08-08
Name/Print	Tod T. Tumey	Registration Number	47.146

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file and by the USPTO to process) an application. Confidentially is governed by 35 U.S. C.12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenar's Office, V.S. Organized to Complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenar's Office, V.S. Organized to Comment of Comment

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that. (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kolfice is to process another examine your submission relation to a patient application or patient. If you do not furnish the requested process another examine your submission relation to the patient application or patient. If you do not furnish the requested the process another examines your submission, which may visually intermediate or for extension or about those when the basic high process another examines your submission, which may visually intermediate or for extension or a submission of the basic high process another examines your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pusuant to 5 U.S.C. 552a(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, cuting an inspection of records concluded by GSAs a part of that apency's responsibility to recommend improvements in records management practices and programs, under suthority of 4d U.S.C. 2004 and 2006. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 122(b) or issuance of a patent pursuant to 35 U.S. C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record via set float in an application which became abandomed or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issuand patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.